This shi s not part of and does not count as a sheet of the ernational application.

PCT For receiving Office	e use only
FEE CALCULATION SHEET	•
Annex to the Request International application No.	· · · · · · · · · · · · · · · · · · ·
Applicant's or agent's file reference 28030-00004  Date stamp of the receiving Office	
Applicant B.C. CHEMICALS LTD. et al.	
CALCULATION OF PRESCRIBED FEES	1
1. TRANSMITTAL FEE	
2. SEARCH FEE	
3. INTERNATIONAL FEE	
Basic Fee The international application contains 23 sheets.	
first 30 sheets	
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remaining sheets additional amount	
Add amounts entered at b1 and b2 and enter total at B	
Designation Fees The international application contains 77 May designations.	,
10 x \$166.00 = \$1660.00 D	
number of designation fees amount of designation fee payable (maximum 10)	·
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4. FEE FOR PRIORITY DOCUMENT (if applicable)	
5. TOTAL FEES PAYABLE	
The designation fees are not paid at this time.	• .
MODE OF PAYMENT	·
authorization to charge deposit account (see below) bank draft coupons  X cheque cash other (specify):  postal money order revenue stamps	
DEPOSIT ACCOUNT AUTHORIZATION (this mode of payment may not be available at all receiving Offices)	
The RO/ is hereby authorized to charge the total fees indicated above to my deposit account.	
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is hereby authorized to charge the fee for preparation and transmittal of the priority document to the International Bureau of WIPO to my deposit account.

Signature

Form PCT/RO/101 (Annex) (January 1999)

Date (day/month/year)

Deposit Account No.

# For receiving Office use only International Application No. REQUEST International Filing Date

international application be processed according to the Patent Cooperation Treaty.	Name of receiving Office and "PCT International Application"
	Applicant's or agent's file reference (if desired) (12 characters maximum) 28030-00004
Box No. I TITLE OF INVENTION	
METHOD FOR THE PREPARATION OF	PHYTOSTEROLS FROM TALL OIL PITCH
Box No. II APPLICANT	
Name and address: (Family name followed by given name; for a designation. The address must include postal code and name of cour address indicated in this Box is the applicant's State (that is, country, of residence is indicated below.)	
B.C. CHEMICALS LTD.	Telephone No.
P.O. Box 6000	Facsimile No.
Prince George, British Columbia	
Canada V2N 2K3	Teleprinter No.
State (that is, country) of nationality:	State (that is, country) of residence:
This person is applicant for the purposes of:  all designated X all designated the United States	States except ates of America
Box No. III FURTHER APPLICANT(S) AND/OR (FURTH	IER) INVENTOR(S)
Name and address: (Family name followed by given name: for a designation. The address must include postal code and name of coun address indicated in this Box is the applicant's State (that is, country) of residence is indicated below.)  WONG, Alfred 3047 West 6th Avenue  Vancouver, British Columbia  Canada V6K 1X4	ry. The country of the of residence if no State  This person is:  applicant only  Applicant and inventor  inventor only (If this check-box is marked, do not fill in below.)
State (that is, country) of nationality:  CA	State (that is, country) of residence:
This person is applicant all designated for the purposes of:	States except
X Further applicants and/or (further) inventors are indicated o	n a continuation sheet.
Box No. IV AGENT OR COMMON REPRESENTATIVE	OR ADDRESS FOR CORRESPONDENCE
The person identified below is hereby/has been appointed to act or of the applicant(s) before the competent International Authorities a	n behalf X agent Common representative
Name and address: (Family name followed by given name; for a designation. The address must include postal co	legal entity, full official de and name of country.)  Telephone No. (604) 687-9444
TURLOCK, Lance A.	
c/o Davis & Company	Facsimile No.
2800 Park Place - 666 Burrard Street	(604) 687-1612
Vancouver, British Columbia	Teleprinter No.
Canada V6C 2Z7	
Address for correspondence: Mark this check-box where n	o agent or common representative is/has been appointed and the
space above is used instead to indicate a special address to w Form PCT/RO/101 (first sheet) (July 1998; reprint January 1999)	hich correspondence should be sent.
i dim i Crivo not (motoneet) (any 1996; tehtin jamary 1999)	See Notes to the request form

Sheet No.	2 .	

Continuation of Box N . III FURTHER APPLICANT(S) A	ND/OR (FURTHER) INVENTOR(S)				
If none of the following sub-boxes is used, this sheet should not be included in the request.					
Name and address: (Family name followed by given name: for a lidesignation. The address must include postal code and name of count address indicated in this Box is the applicant's State (that is, country) of residence is indicated below.)  NORMAN, Hugh S. O.  864 Reid Crescent  Prince George, British Columbia  Canada V2N 3W8	egal entity, full official ry. The country of the of residence if no State  This person is:  applicant only  X applicant and inventor  inventor only (If this check-box is marked, do not fill in below.)				
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Name and address: (Family name followed by given name; for a ladesignation. The address must include postal code and name of count address indicated in this Box is the applicant's State (that is, country) of residence is indicated below.)  MACMILLAN, Angus Kirke  13 Sherwood Place  Tsawwassen, British Columbia  Canada V4L 2C7	ry. The country of the of residence if no State  This person is:  applicant only  applicant and inventor  inventor only (If this check-box is marked, do not fill in below.)				
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Further applicants and/or (further) inventors are indicated or	n another continuation sheet.				

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Box I	No.V	DESIGNATION OF STATES			· · · · · · · · · · · · · · · · · · ·			
The	The following designations are hereby made under Rule 4.9(a) (mark the applicable check-boxes; at least one must be marked):							
	Regional Patent							
X	<u> </u>							
	AF	ZW Zimbabwe, and any other State which is a Control						
X	EA	Eurasian Patent: AM Armenia, AZ Azerbaijan, BY Belarus, KG Kyrgyzstan, KZ Kazakhstan, MD Republic of Moldova, RU Russian Federation, TJ Tajikistan, TM Turkmenistan, and any other State which is a Contracting State						
		of the Eurasian Patent Convention and of the PCT						
Ø	EP	DK Denmark, ES Spain, FI Finland, FR France, GB MC Monaco, NL Netherlands, PT Portugal, SE Sweet Patent Convention and of the PCT	European Patent: AT Austria, BE Belgium, CH and LI Switzerland and Liechtenstein, CY Cyprus, DE Germany, DK Denmark, ES Spain, FI Finland, FR France, GB United Kingdom, GR Greece, IE Ireland, IT Italy, LU Luxembourg, MC Monaco, NL Netherlands, PT Portugal, SE Sweden, and any other State which is a Contracting State of the European Patent Convention and of the PCT					
X	OA	GA Gabon, GN Guinea, GW Guinea-Bissau, ML Ma	li, MI and	RMau a Cor	Republic, CG Congo, CI Côte d'Ivoire, CM Cameroon, uritania, NE Niger, SN Senegal, TD Chad, TG Togo, and stracting State of the PCT (if other kind of protection or treatment			
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Precautionary Designation Statement: In addition to the designations made above, the applicant also makes under Rule 4.9(b) all other designations which would be permitted under the PCT except any designation(s) indicated in the Supplemental Box as being excluded from the scope of this statement. The applicant declares that those additional designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be regarded as withdrawn by the applicant at the expiration of that time limit (Confirmation of a designation consists of the filing of a notice specifying that designation and the payment of the designation and confirmation fees. Confirmation must reach the receiving Office within the 15-month time limit.)

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Box No. VI PRIORITY C	LAIM		Further prio	rity claims are indicated	in the Supplemental Box.	
Filing date Number of earlier application of earlier application			Where earlier application is:			
of earlier application (day/month/year)	of earlier ap	pplication	national application:		international application:	
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From the INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

TURLOCK, Lance A. Davis & Company 2800 Park Place 666 Burrard Street Vancouver, B.C. V6C 2Z7 **CANADA** 

NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL PRELIMINARY **EXAMINATION REPORT** 

(PCT Rule 71.1)

Date of mailing (day/month/year)

02.06,2000

Applicant's or agent's file reference 28030-00004

IMPORTANT NOTIFICATION

International application No. PCT/CA99/00150

International filing date (day/month/year) 19/02/1999

Priority date (day/month/year) 20/02/1998

**Applicant** 

B.C. CHEMICALS LTD. et al.

- 1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
- 2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
- 3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

#### 4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/

Authorized officer

Brell, S

European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465

Tel.+49 89 2399-7271



## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/CA99/00150

<ol> <li>Basis of the repo</li> </ol>	r	l
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1.	This report has been drawn on the basis of (substitute sheets which have been furnished to the receiving Office in
	response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to
	the report since they do not contain amendments.):

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	1,2	5-13	as originally filed		. •	
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### PATENT COOPERATION TREATY

### PCT

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicants	01 20	ent's file reference	<del></del>				
1 ''	11		FOR FURTHER AC	TION		ation of Transmittal of International y Examination Report (Form PCT/IPEA/	/416)
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and i	s tran:	smitted to the applicant	according to Article 36.	,		ernational Preliminary Examining A	uthority
2. This	REPC	ORT consists of a total or	f 4 sheets, including this	cover s	heet.		
t (	This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).  These annexes consist of a total of 7 sheets.						
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### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/CA99/00150

V. Reasoned stat ment und r Article 35(2) with r gard to novelty, inventive step r industrial applicability; citations and xplanations supporting such statement

1. Statement

Novelty (N)

Yes:

Claims 1-25

No:

Claims 26

Inventive step (IS)

Yes:

Claims 1-25

No:

Claims 26

Industrial applicability (IA)

Yes: No: Claims 1-26 Claims

2. Citations and explanations

see separate sheet

#### INTERNATIONAL PRELIMINARY **EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/CA99/00150

To section V:

Novelty:

MALIK, LUBOMIR ET AL: 'Isolation of phytosterols from tall - oil rosin' describes a process for the isolation of phytosterols, said process involving four distillation steps which are carried out in recombinant parallel paths (e.g.the light fraction of the second distillation is combined with the bottom fraction of a third distillation, and that combined fractions are re-distilled).

The present process is distinguished from said process in that it involves a sequence of distillation steps which are carried out in series, and in that the fractions are not mixed prior to further treatment.

Thus the processes as defined in claims 1-25 are novel in view of the prior art process.

Claim 26: Phytosterols are well known in the art. Claim 26 does not contain any technical feature that would allow to distinguish the claimed compositions from the phytosterols described in the prior art. The fact that the phytosterols are obtained by a novel process does not bring about novelty (or inventive step) of the phytosterols per se.

#### Inventive step:

Closest prior art appears to be the process described in Malik et al. referred to above. Said prior art process results in a recovery of phytosterols of 80-90% by weight. The present process is advantageous over the aforementioned pror art process since it avoids parallel distillations and uses fewer distillation steps, whereas the percentage of recovered phytosterols is equally high (about 90%, cf. table 2). Such improvement is not deducible from the prior art, an inventive step can therefore be acknowledged.



In U.S. Patent No. 5,097,012 granted on 17 March 1992, Thies et al. disclose a method for the isolation of sterols from crude tall oil by water extraction at elevated temperatures and pressures.

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In U.S. Patent No. 3,943,117 granted on 9 March 1976, Force discloses a process for saponifying tall oil pitch in which a water-soluble cationic amine is used in conjunction with an alkali. In U.S. Patent No. 4,524,024 granted on 18 June 1985, Hughes teaches the hydrolysis of tall oil pitch at elevated temperatures to increase the recovery of fatty acids from tall oil pitch. In U.S. Patent No. 3,887,537 granted on 3 June 1975, Harada et al. disclose the recovery of fatty acids and rosin acids from tall oil pitch by first saponifying tall oil pitch with an alkali metal base and a low molecular weight alcohol, and then introducing the reacted mixture into a thin film evaporator to remove low-boiling matter such as water, alcohol use and light unsaponifiables. The bottom fraction from the first evaporator is next fed to a second thin film evaporator in which the unsaponifiables including sterols are removed as the light ends and a molten soap is recovered as the bottom fraction. Fatty acids and rosin acids are recovered from the molten soap fraction by acidulation conventionally with a mineral acid. In U.S. Patent No. 3,926,936 granted on 16 December 1975, Lehtinen teaches the recovery of fatty acids and rosin acids from tall oil pitch by reacting tall oil pitch with an alkali at 200 to 300 degrees Celsius, in the amount of 5 to 25% of tall oil pitch, prior to vacuum distillation of the heated mixture to recover the fatty acids and rosin acids in the distillate fraction.

Reference is also made to Chemical Abstracts, vol. 112, no. 20, 14 May 1990, Columbus, Ohio US; abstract no. 181758, MALIK, Lubomir et al: "Isolation of phytosterols from tall - oil rosin", XP002104877 & CS 256 092 A (Czech). Malik et al. disclose a process for extracting phytosterols which includes the use of four distillation stages. Product flow is split into parallel distillation paths in a first distillation stage then, following further distillation in each parallel path, is partially recombined prior to a final distillation stage. To achieve high purity phytosterols, the output from the final distillation



stage is subjected to two stages of crystallization utilizing relatively large amounts of solvent.

#### SUMMARY OF THE INVENTION

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In a broad aspect of the present invention there is provided a new and improved method of preparing phytosterols from tall oil pitch containing steryl esters, the method comprising the steps of:

- (a) converting the steryl esters to free phytosterols while in the pitch to produce a modified pitch containing the free phytosterols;
- (b) distilling the modified pitch in a first evaporator to remove light ends from the modified pitch and produce a bottom fraction containing the free phytosterols;
- (c) distilling only the bottom fraction in a second evaporator to produce a light phase distillate containing the free phytosterols;
- (d) dissolving only the light phase distillate in a solvent comprising an alcohol to produce a solution containing the free phytosterols;
- (e) cooling the solution to produce a slurry with the free phytosterols crystallized in the slurry; and,
  - (f) washing and filtering the slurry to isolate the crystallized phytosterols.

Preferably, the step of converting the steryl esters to free phytosterols comprises the steps of saponifying the tall oil pitch with an alkali metal base, neutralizing the saponified pitch with an acid, and heating the neutralized pitch to remove water. The resulting pitch with such water removed defines the modified pitch.

Unlike the process of Malik et al., the foregoing process enables the preparation of high purity phytosterol crystals from tall oil pitch with only two distillation stages and only one stage of crystallization, and to do so with the use of a comparatively small amount of solvent. Nevertheless, it may be considered desirable in some cases to achieve phytosterol yields with even higher crystal purity. In accordance with another embodiment of the invention, a marginal improvement is achieved as follows:



- (a) producing a light phase distillate containing free phytosterols in the manner described in steps (a) to (c) above;
- (b) re-distilling only the light phase distillate so produced to enhance the concentration of free phytosterols in the light phase distillate;
- (c) dissolving only the re-distilled light phase distillate in a solvent comprising an alcohol to produce a solution containing the free phytosterols; and,
- (d) continuing the procedure as in steps (d) and (f) above to isolate crystallized phytosterols.

Although this procedure involves additional distillation steps, the amount of
alcohol required during the crystallization stage remains small compared to the case of
Malik et al.

#### BRIEF DESCRIPTION OF THE DRAWINGS

The FIGURE shows a schematic flow diagram for the preparation of high purity phytosterol crystals from tall oil pitch in accordance with the present invention.

#### DESCRIPTION OF PREFERRED EMBODIMENT

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In accordance with the present invention, the isolation of phytosterols from tall oil pitch first requires converting steryl esters present in the pitch to free phytosterols while in the pitch. The result is a modified pitch containing free phytosterols.

It is contemplated that the required conversion may be accomplished by various methods. In the FIGURE, the conversion step is indicated by block 30 (shown in broken outline) which receives an incoming feed of tall oil pitch 1 and produces modified pitch 11 as an output. The presently preferred method of conversion involves the use of an alkali base treatment and is indicated by the elements contained within block 30.

PCT/CA99/00150

#### WE CLAIM:

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15.

- 1. A method of preparing phytosterols from tall oil pitch (1) containing steryl esters, said method comprising the steps of:
  - (a) converting said steryl esters to free phytosterols while in said pitch to produce a modified pitch (11) containing said free phytosterols;
  - (b) distilling said modified pitch (11) in a first evaporator (12) to remove light ends (13) from said modified pitch and produce a bottom fraction (14) containing said free phytosterols;
  - (c) distilling only said bottom fraction (14) in a second evaporator (15) to produce a light phase distillate (16) containing said free phytosterols;
  - (d) dissolving only said light phase distillate (16) in a solvent (21) comprising an alcohol to produce a solution containing said free phytosterols;
  - (e) cooling said solution to produce a slurry (19) with said free phytosterols crystallized in said slurry; and,
  - (f) washing and filtering said slurry (19) to isolate said crystallized phytosterols (22).
- 2. A method as defined in claim 1, wherein said modified pitch (11) comprises less than 1% water by weight.
- 3. A method as defined in claim 1 or 2, wherein said solvent (21) comprises a low molecular weight monohydric alcohol.
- 4. A method as defined in claim 1 or 2, wherein said solvent (21) comprises a low molecular weight monohydric alcohol and water.
- 5. A method as defined in claim 1 or 2, wherein said slurry (19) is washed and filtered using a solvent like said solvent (21) used to dissolve said light phase distillate.
- 25 6. A method as defined in claim 1, wherein said step of converting said steryl esters to free phytosterols comprises the steps of:
  - (a) saponifying said tall oil pitch (1) with an alkali metal base (2);

- (b) neutralizing said saponified pitch with an acid (5); and,
- (c) heating said neutralized pitch to remove water, the resulting pitch with such water removed defining said modified pitch (11).
- 7. A method as defined in claim 6, wherein said alkali metal base (2) is selected from the group consisting of:
  - (a) sodium hydroxide;
  - (b) potassium hydroxide;
  - (c) sodium hydroxide and potassium hydroxide:
- 8. A method as defined in claim 7, wherein in the weight percentage of alkali metal base (2) to tall oil pitch (1) is in the range of 1% to 15%.
  - 9. A method as defined in claim 7, wherein said saponification is conducted at a temperature in the range of 100 to 250 deg. C for a period in the range of 60 to 300 minutes.
  - 10. A method as defined in claim 6, wherein said acid (5) is an organic acid.
- 15 11. A method as defined in claim 6, wherein said acid (5) is a mineral acid.
  - 12. A method as defined in claim 11, wherein said mineral acid (5) is selected from the group consisting of:
    - (a) sulphuric acid;
    - (b) hydrochloric acid;
- 20 (c) phosphoric acid;
  - (d) a combination of acids comprising two or more of sulphuric acid, hydrochloric acid and phosphoric acid.
  - 13. A method as defined in claim 6, wherein said neutralization is conducted at a temperature in the range of 10 to 100 deg. C for a period in the range of 1 to 10 hours.



- 14. A method as defined in claim 6, wherein said neutralized pitch has a water phase pH in the range of 4 to 7.
- 15. A method as defined in claim 6, wherein said heating step comprises heating at a temperature in the range 90 to 100 deg. C for a time sufficient to effect the bulk disengagement of water from the organic phase.

- 16. A method as defined in claim 15, wherein said heating step further comprises heating under vacuum conditions such that said modified pitch (11) comprises less than 1% water by weight.
- 17. A method as defined in claim 1 or 6, wherein said light ends are removed in a
  wiped film evaporator (12) operating at a pressure in the range of 0.1 to 10 millibars and at
  a temperature in the range 160 to 280 deg. C.
  - 18. A method as defined in claim 1 or 6, wherein said bottom fraction is evaporated in a wiped film evaporator (15) operating at a pressure in the range of 0.01 to 1.0 millibars and at a temperature in the range 180 to 300 deg. C.
- 15 19. A method as defined in claim 6, wherein said solvent (21) comprises a low molecular weight monohydric alcohol.
  - 20. A method as defined in claim 6, wherein said solvent (21) comprises a low molecular weight monohydric alcohol and water.
- 21. A method as defined in claim 1 or 6 in which the crystallization of phytosterols is effected at a temperature in the range of 0 to 35 deg. C.
  - 22. A method of preparing phytosterols from tall oil pitch (1) containing steryl esters, said method comprising the steps of:
    - (a) converting said steryl esters to free phytosterols while in said pitch to produce a modified pitch (11) containing said free phytosterols;



- (b) distilling said modified pitch (11) in a first evaporator (12) to remove light ends (13) from said modified pitch and produce a bottom fraction (14) containing said free phytosterols;
- (c) distilling only said bottom fraction (14) in a second evaporator (15) to produce a light phase distillate (16) containing said free phytosterols;
- (d) re-distilling only said light phase distillate (16) to enhance the concentration of free phytosterols in said light phase distillate;
- (e) dissolving only said re-distilled light phase distillate in a solvent (21) comprising an alcohol to produce a solution containing said free phytosterols;
- (f) cooling said solution to produce a slurry (19) with said free phytosterols crystallized in said slurry; and,
- (g) washing and filtering said slurry (19) to isolate said crystallized phytosterols (22).
- 23. A method as defined in claim 22, wherein said solvent (21) further comprises water added in a proportion up to 35% by weight relative to the organic solvent phase.
  - 24. A method as defined in claim 23, wherein the weight ratio of solvent to distillate is between 0.3 to 2.0.
  - 25. A process according to claim 19, 20 or 24 in which the alcohol is selected from:
    - (a) methanol;
- (b) ethanol;

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- (c) 2-propanol;
- (d) a combination of alcohols comprising two or more of methanol, ethanol and2-propanol.
- 26. Phytosterols prepared from tall oil pitch in accordance with the method as definedin any one or more of the preceding claims.

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#### INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	(Form PCT/ISA/	of Transmittal of International Search Report 220) as well as, where applicable, item 5 below.
28030-00004	ACTION	
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)
PCT/CA 99/00150	19/02/1999	20/02/1998
Applicant		·
B.C. CHEMICALS LTD. et al	•	
This International Search Report has been according to Article 18. A copy is being tra	n prepared by this International Searching Au ansmitted to the International Bureau.	thority and is transmitted to the applicant
This later should Count Depart services	of a hard of 2 about	
This International Search Report consists  It is also accompanied by	of a total of sheets.  a copy of each prior art document cited in this	s report.
Basis of the report		<del></del>
·	· · · · international search was carried out on the ba	usis of the international application in the
language in which it was filed, unl	ess otherwise indicated under this item.	on the mornauonal application in the
the international search w Authority (Rule 23.1(b)).	as carried out on the basis of a translation of	the international application furnished to this
b. With regard to any nucleotide an	d/or amino acid sequence disclosed in the i	ntemational application, the international search
was carried out on the basis of the contained in the internation	e sequence listing : nal application in written form.	
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furnished subsequently to	this Authority in written form.	
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the statement that the sub international application as	sequently turnished written sequence listing of sequence listing of the sequen	does not go beyond the disclosure in the
the statement that the info furnished	rmation recorded in computer readable form	is identical to the written sequence listing has been
2. Certain claims were four	nd unsearchable (See Box I).	•
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X the text is approved as su	bmitted by the applicant.	
the text has been establish	hed by this Authority to read as follows:	
5. With regard to the abstract,		•
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within one month from the	ned, according to Hule 38.2(b), by this Author date of mailing of this international search re	nty as it appears in Box III. The applicant may, sport, submit comments to this Authority.
6. The figure of the drawings to be publi	ished with the abstract is Figure No.	1
as suggested by the applie	cant.	None of the figures.
X because the applicant faile	ed to suggest a figure.	
	characterizes the invention.	

According to International Patent Classification (IPC) or to both national classification and IPC

#### B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) IPC 6 C07J C11B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DUCUM	NTS CONSIDERED TO BE RELEVANT	
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	CHEMICAL ABSTRACTS, vol. 112, no. 20, 14 May 1990 Columbus, Ohio, US; abstract no. 181758, MALIK, LUBOMIR ET AL: "Isolation of phytosterols from tall - oil rosin" page 139; column 1;	1-25
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Y Further documents are listed in the continuation of box C.	Patent family members are listed in annex.
*Special categories of cited documents:  "A" document defining the general state of the art which is not considered to be of particular relevance  "E" earlier document but published on or after the international filing date  "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)  "O" document referring to an oral disclosure, use, exhibition or other means  "P" document published prior to the international filing date but later than the priority date claimed	<ul> <li>'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</li> <li>'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</li> <li>'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</li> <li>'&amp;' document member of the same patent family</li> </ul>
Date of the actual completion of the international search  4 June 1999	Date of mailing of the international search report 25/06/1999
Name and mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2  NL - 2280 HV Rijswijk  Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  Fax: (+31-70) 340-3016	Authorized officer Watchorn, P

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